

CLAIMS

- 5 1. A peptide containing at least 6 amino acid residues and having at least 70% homology with part or all of the sequence
AEFHRWSSYMVHWK.
2. A peptide comprising or consisting of the sequence YMVH or MVHW or VHWK and having at least 70% homology with part or all of the
10 sequence
AEFHRWSSYMVHWK.
- 3 A mixture of the peptide of claim 1 or claim 2 with another peptide having at least 4 amino acid residues and having at least 70% homology with the β -amyloid precursor sequence
15 DAEFRHDSGYEVHHQK.
4. A probe consisting of the peptide of claim 1 or claim 2 or the mixture of claim 3, labelled with a signal moiety, or immobilised on a support.
5. A compound which competes with the peptide of claim 1 or
20 claim 2 for binding to a receptor therefor and which thereby inhibits the biological activity of the said peptide.
6. A compound as claimed in claim 5, wherein the biological activity is modulating a calcium-channel-opening activity.
7. A compound as claimed in claim 5 or claim 6, which is
25 capable of crossing the blood-brain barrier.
8. An antibody to the peptide of claim 1 or claim 2.
9. An antibody as claimed in claim 8 which is of the IgG class.
10. An antibody fragment or chimeric or humanised antibody comprising variable regions of the antibody of claim 8 or claim 9.

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11. A method of preparing a composition for treatment of disorders of the central nervous system or stroke or cancer, which method comprises bringing a compound according to any one of claims 5 to 10 into a form for human administration.
- 5 12. A method of preparing a composition for controlling cytoplasmic calcium ion concentration *in vivo*, which method comprises bringing a compound according to any one of claims 5 to 10 into a form for human administration.

ADD
A1

Add D2